Intestinal circulation, oxygenation and metabolism is not affected by oleic acid lung injury

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Summary

This study was performed to establish a platform for further studies on effects of ventilatory treatment modalities on the intestines during mechanical ventilation of acute lung injury (ALI). We tested the hypotheses that oleic acid (OA) infusion causes changes in intestinal circulation, oxygenation and metabolism, and that OA is distributed to tissues outside the lung. This was performed as an experimental, prospective and controlled study in an university animal research laboratory. Thirteen juvenile anaesthetized pigs were used in the main study, where seven were given an intravenous infusion of 0.1 ml kg⁻¹ OA and six served as control (surgery only). In a separate study, four animals were given an intravenous infusion of 0.1 ml kg⁻¹ ³H-labelled OA. We measured systemic and mesenteric (portal venous blood flow, jejunal mucosal perfusion) haemodynamic parameters, mesenteric oxygenation (jejunal tissue oxygen tension) and systemic cytokines (tumour necrosis factor-α and interleukin-6). We calculated mesenteric lactate flux and mesenteric oxygen delivery, uptake and extraction ratio. In the animals given ³H-OA, we measured ³H-OA in different tissues (lungs, heart, liver, kidney, stomach, jejunum, colon and arterial blood). We found that OA given intravenously is distributed in small amounts to the intestines. This intestinal exposure to OA does not cause intestinal injury when evaluating mesenteric blood flow, metabolism or oxygenation. OA infusion induced a moderate increase in mean pulmonary arterial pressure and a decrease in PaO₂/Fraction inspired O₂ (P/F) ratio, giving evidence of severe lung injury. Consequently, the OA lung injury model is suitable for studies on intestinal effects of ventilatory treatment modalities during mechanical ventilation of ALI.

Introduction

The intestine, suggested to be a key organ in the development of multiple organ failure (Mainous et al., 1995; Gullo & Berlot, 1996; Nieuwenhuijzen & Goris, 1999), is well known to be negatively affected by ventilator treatment both in healthy and injured lungs (Winsø et al., 1986; Claesson et al., 2003; Guery et al., 2003; Imai et al., 2003; Lehtipalo et al., 2003; Plotz et al., 2004). Research from our own group has shown that treatment with positive end-expiratory pressure (PEEP) reduces portal blood flow during general surgery in man (Winso et al., 1986). Further, we have shown that, in a porcine model, treatment with modest levels of PEEP causes ischaemia when intestinal perfusion pressure is reduced (Lehtipalo et al., 2003). We have also shown that lung recruitment manoeuvres in patients with acute lung injury (ALI) decreases gastric mucosal blood flow (Claesson et al., 2003). Also, in lung injury, the resulting general hypoxia may cause redistribution of regional perfusion, disfavouring splanchnic oxygen delivery.

Others have shown (Plotz et al., 2004) that the intestines might be affected negatively by several other mechanisms during mechanical ventilation. Systemically released cytokines and apoptotic mediators from the lung might cause increased intestinal permeability and intestinal epithelial apoptosis promoting bacterial translocation (Guery et al., 2003; Imai et al., 2003). From this review, it is obvious that many factors affect the intestines during mechanical ventilation and that the interactions between these factors are complex. It is important to elucidate these interactions as there are experimental evidence gathering that there might be clinically relevant interactions between mechanical ventilation in lung injury, changes in intestinal permeability and subsequent development of multiple organ failure (Doig et al., 1998; Dreyfuss & Saumon, 1998).
The main purpose of this study was to establish a platform for further studies on effects of different ventilatory treatment modalities on the intestines during mechanical ventilation of ALI. The oleic acid (OA) lung injury model is one of the accepted standards to study ventilatory management in ALI (Kloot et al., 2000; Neumann & Hedenstierna, 2001; Pelosi et al., 2001; Lim et al., 2004).

However, it remains unclear if OA is solely distributed to the lung, or whether it also can be found in other organs. If the latter were the case, possible local effects and, in particular, further regional changes in organ perfusion and metabolism might occur. This in turn might result in direct impairment of organ function other than in the lung alone. Finally, only scarce data exist, documenting stability of the OA model for several hours.

First, in the main study, we aimed to test the hypothesis that OA infusion has intrinsic effects on intestinal circulation, metabolism and oxygenation. Further, in a separate study, we aimed to test the hypothesis that infused OA would be distributed to tissues outside the lung. To this end, we used an established experimental model of OA-induced ALI.

**Methods**

The study was approved by the University Animal Experiment Ethics Committee. Thirteen female juvenile pigs, with a mean weight of 30·8 ± 1·1 kg, were used in the main study. Another four pigs, with a mean weight of 27·7 ± 0·8 kg, were used in a separate study of tissue distribution of tritium (3H)-labelled OA. All procedures were carried out according to the guidelines of the National Institute of Health Guide for the care and use of laboratory animals (7th edn, 1996).

**Anaesthesia**

Animals were fasted overnight with free access to water. After premedication with ketamine, azaperon and atropine i.m., anaesthesia was induced by sodium pentobarbital (10 mg kg⁻¹, i.v.) followed by continuous infusions of sodium pentobarbital 4–6 mg kg⁻¹ h⁻¹ i.v., midazolam 0·3 mg kg⁻¹ h⁻¹ i.v. and fentanyl 20 µg kg⁻¹ h⁻¹ i.v. No muscle relaxants were used. After tracheostomy, mechanical ventilation with oxygen in air (30% O₂) was performed using pressure-regulated volume control (Evita 4, Dräger, Germany), tidal volume of 10 ml kg⁻¹ and frequency of 20–30 b min⁻¹. Ventilation was adjusted to normocapnia as judged by end-tidal CO₂ levels (Artela; Artema Medical AB, Sundbyberg, Sweden) and arterial blood gas analyses (ABL-5 autoanlyser; Radiometer, Copenhagen, Denmark). Blood samples for lactate concentration were analysed by an automated analyser (YSI Sport 2300 Stat Plus; Yellow Springs Instruments, Inc., Yellow Springs, Ohio, USA). All animals received i.v. infusions of Ringer’s acetate 10 ml kg⁻¹ h⁻¹ and hydroxyethylstarch 200/0.6 (Voluven, Fresenius Kabi, Sweden) 2 ml kg⁻¹ h⁻¹, with additional boluses given to keep central venous pressure (CVP) in the range of 5–7 mmHg during the experiment. Core temperature was kept between 37 and 39°C using heating blankets.

**Instrumentation and measurements**

Instrumentation of the animals was performed as previously described regarding intravascular catheters, portal blood flow (Qport), partial tissue O₂ pressure (PtiO₂) and jejunal mucosal perfusion (JMP) measurements (Lehtipalo et al., 2003). In eight pigs a catheter (Pulsiocath 4F; Pulsion Medical Systems AG, Munich, Germany) was inserted via a branch of the left carotid artery for monitoring of pulse contour cardiac output (CO). The pulse contour computer continuously calculated the arterial pulse contour CO. Calibration of the system was performed twice during the experiment according to the manufacturer’s instructions by determination of the arterial thermodilution CO, using three to four injections of 20 ml iced 0.9% saline (Sakka et al., 1999). In those pigs where continuous CO monitoring was not available, CO was measured using the pulmonary artery catheter. Thermodilution CO data (5 ml of iced 0.9% NaCl as indicator) are presented as means of three consecutive measurements within 2 min, not differing more than 10%. Splenectomy was performed in all animals to eliminate dilution of portal venous blood. All pressure transducers (System DPT-6000, PVB; Triplus, Kungsbacka, Sweden) were calibrated to atmospheric pressure at the level of the right atrium by a saline column. Continuous data were acquired with Acknowledge® III (Biopac Systems Inc., Goleta, California, USA) as previously described (Lehtipalo et al., 2003).

**Cytokine measurements**

Arterial blood was centrifuged at 4°C at 915 g for 20 min and serum was collected after centrifugation. The serum samples were then frozen at −70°C awaiting analysis. Cytokine ELISA kits for tumour necrosis factor (TNF)-α and interleukin (IL)-6 were purchased from R&D Systems Europe Ltd (Abingdon, UK). Assays were performed as recommended by the manufacturer with some minor modifications. Briefly, capture antibodies were incubated in microtitre 96-well ELISA plates (Nunc-Immuno Plate with Max Sorb surface, Tamro MedLab AB, Mölndal, Sweden) overnight at room temperature. Plates were then washed (0.05% Tween in physiologically buffered saline) and non-specific binding sites in each well were blocked by incubating with a buffer consisting of bovine serum albumin (1%) and sucrose (5%) in physiologically buffered saline. One hour later, 100 µl of each serum sample was added to the wells. Serial dilutions of serum samples were performed in order to keep optical density values within the linear range of the twofold serial dilution standard curve. Final analysis was performed with serum dilutions 1:10. After 2 h incubation, bound cytokines were detected using a biotin-coupled detection antibody which was quantified using a strepavidin-peroxidase conjugate. After repeated washing, a ready-to-use peroxidase substrate with 3,3′,5,5′-tetramethylbenzidine and hydrogen peroxide (Sigma-
Aldrich, Stockholm, Sweden) was used. The plates were then incubated at room temperature, allowing the reaction to proceed to the desired extinction. The reaction was terminated by adding 50 μl of 1 M H2SO4 to each well. The soluble product was read at 450 nm in a Labsystems iEMS ELISA reader.

**Determination of tissue 3H oleic acid radioactivity**

The individual organs were harvested and placed in a sink, and blood was allowed to drain freely. Thereafter, three 1-g tissue samples from each organ (heart, lungs, liver, kidneys, stomach, jejunum and colon) were separately homogenized with a tissue homogenisator (Polytron; Kinematica, Basel, Switzerland) and extracted in 30 ml chloroform–methanol 2:1 per gram and then left overnight. One millilitre of arterial blood was also sampled and treated accordingly. An aliquot of the extract was centrifuged for 10 min to remove tissue residues. Three millilitre of the clear phase was added to 1-2 ml 0·1 M HCl and shaken. After an additional centrifugation to separate the two phases, the upper phase and protein layer were discarded, and 1 ml of the lower phase was transferred to a scintillation vial. After the chloroform was evaporated over night, scintillation fluid (Optiphase HiSafe III; Perkin-Elmer, Stockholm, Sweden) was added and allowed to mix for another 24 h, after which the sample activity was counted in a Wallace 1414 WinSpectral β-counter. Estimation of the contribution of 3H-OA radioactivity from residual blood was performed based on previously reported individual organ residual blood volume (Hansard, 1956).

**Experimental protocol, main study**

After completed instrumentation the animals were allowed to stabilize for 60 min and the OA group was divided into two groups, a control group (n = 6) and an OA group (n = 7). In the OA group, lung injury was established by a slow 30-min intravenous infusion of 0·1 ml kg–1 OA (Apoteksbolaget, Göteborg, Sweden) suspended in 20 ml saline via the central venous catheter. Inspired oxygen concentration was adjusted between a FiO2 of 0·3 to 1·0 to maintain oxygen saturation above 92%. During the OA infusion ventilation was increased when needed to maintain normoventilation. Blood pressure was stabilized with additional colloid infusion (500–1000 ml to maintain CVP at 5–7 mmHg) and repeated doses of adrenaline 5–20 μg (to a total of 100–250 μg). After the 30-min OA infusion, PEEP 5 cm H2O was added and the animals were allowed to recover for 80 min.

In the control group only PEEP 5 cm H2O was added after 30 min. No extra fluids or adrenaline was given. In both groups, data sets were recorded at time zero and 120, 150, 175, 200 and 240 min after time zero.

Recordings included heart rate (HR), mean arterial pressure (MAP), CVP, mean pulmonary arterial pressure (MPAP), CO, QPORT, portal venous pressure (PvORT), PetO2, and JMP. Blood samples for blood gas analyses and oxygen saturation measurements were drawn from aortic and portal venous catheters in conjunction with the haemodynamic recordings. At time zero, 120 and 240 min thereafter mixed venous blood samples for blood gas analyses were collected, venous admixture (Qs/Qt) was calculated and cytokines were measured. Animals were killed with an intravenous infusion of KCl under deep barbiturate anaesthesia at the end of the experiment.

**Experimental protocol, 3H-OA distribution**

In a separate group of four animals, the distribution of OA to different tissues was studied. These animals were anaesthetized and instrumented as described above, with the exception that the only intra-abdominal instrumentation performed was splenectomy.

After completed instrumentation the animals were allowed to stabilize for 30 min. 3H-labelled OA 0·1 ml kg–1 (200 μCi) was then injected as a slow i.v. infusion during 30 min in the central venous catheter as described above. The animals received the same treatment regarding ventilation, fluids and adrenaline as the animals in the OA group in the main study. The animals were followed for another 45 min after the OA infusion. In this distribution study, HR, MAP, CVP and MPAP were continuously recorded. Arterial and mixed venous blood gases were sampled at 15 min intervals. All animals were killed with an intravenous infusion of KCl under deep barbiturate anaesthesia at the end of the experiment. After death, tissue samples (three 1-g samples from each tissue) and 1 ml arterial blood were taken for determination of 3H-labelled OA radioactivity.

**Statistics**

All values are given as mean ± SEM. Comparisons were made between groups at time-points zero, 120 and 240 min. Comparisons were made within groups between time-point zero and 120 min and between 120 and 240 min. For comparisons between groups an independent samples t-test was used. For comparisons within groups a paired samples t-test was used. A P<0·05 was considered significant for all tests. The statistical analysis was performed with the SPSS software package (version 11·5; SPSS Inc., Chicago IL, USA).

**Results, main study**

The results are reported as changes occurring during the first 120 min after time zero (acute phase) followed by changes occurring between 120 and 240 min after time zero (late phase). At time zero there were no significant differences between groups apart from slightly higher pH in the OA group.

**Acute phase, changes within groups**

In the control group MAP and QPORT decreased (QPORT from 1059 ± 153 to 800 ± 103 ml min–1). CO decreased although not significantly (Table 1, Fig. 1). Mesenteric oxygen delivery
(M-DO₂) decreased from 132 ± 18 to 104 ± 14 ml min⁻¹ while mesenteric oxygen extraction ratio (MO₂-ER) increased from 21 ± 3% to 28 ± 3%.

In the OA group MAP, CO, QPORT, pH and P/F ratio decreased (QPORT from 900 ± 75 to 681 ± 47 ml min⁻¹) while MPAP and PaCO₂ increased (Table 1, Fig. 1). M-DO₂ decreased from 110 ± 11 to 82 ± 7 ml min⁻¹ while MO₂-ER increased from 25 ± 2% to 36 ± 1%.

Table 1: Circulatory and respiratory parameters.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Time zero</th>
<th>120 min</th>
<th>240 min</th>
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<tr>
<td>MAP (mmHg)</td>
<td>OA 98 ± 4</td>
<td>85 ± 4*</td>
<td>80 ± 4&quot;</td>
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<tr>
<td></td>
<td>Control 92 ± 2</td>
<td>82 ± 4*</td>
<td>80 ± 3</td>
</tr>
<tr>
<td>CVP (mmHg)</td>
<td>OA 6 ± 0</td>
<td>8 ± 0</td>
<td>6 ± 0*</td>
</tr>
<tr>
<td></td>
<td>Control 6 ± 0</td>
<td>7 ± 1</td>
<td>8 ± 1</td>
</tr>
<tr>
<td>MPAP (mmHg)</td>
<td>OA 22 ± 1</td>
<td>37 ± 2*</td>
<td>30 ± 2&quot;</td>
</tr>
<tr>
<td></td>
<td>Control 23 ± 1</td>
<td>22 ± 1</td>
<td>25 ± 3</td>
</tr>
<tr>
<td>CO (L min⁻¹)</td>
<td>OA 4-8 ± 0-3</td>
<td>3-8 ± 0-2*</td>
<td>3-7 ± 0-2</td>
</tr>
<tr>
<td></td>
<td>Control 4-7 ± 0-5</td>
<td>3-5 ± 0-3</td>
<td>3-5 ± 0-1</td>
</tr>
<tr>
<td>PH</td>
<td>OA 7-54 ± 0-01§</td>
<td>7-40 ± 0-02*§</td>
<td>7-44 ± 0-04</td>
</tr>
<tr>
<td></td>
<td>Control 7-48 ± 0-01</td>
<td>7-48 ± 0-01</td>
<td>7-47 ± 0-01</td>
</tr>
<tr>
<td>PaCO₂ (kPa)</td>
<td>OA 4-5 ± 0-3</td>
<td>5-9 ± 0-3*</td>
<td>5-6 ± 0-7</td>
</tr>
<tr>
<td></td>
<td>Control 5-0 ± 0-1</td>
<td>4-9 ± 0-1</td>
<td>5-0 ± 0-2</td>
</tr>
<tr>
<td>PaO₂ (kPa)</td>
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<td>15-7 ± 2-0</td>
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<td>Control 14-5 ± 0-5</td>
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<td>13-8 ± 0-8&quot;</td>
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<tr>
<td>P/F (kPa)</td>
<td>OA 49-2 ± 4-0</td>
<td>27-6 ± 4-8*§</td>
<td>25-9 ± 4-2*§</td>
</tr>
<tr>
<td></td>
<td>Control 48-3 ± 1-5</td>
<td>51-3 ± 1-4</td>
<td>46-2 ± 2-6&quot;</td>
</tr>
<tr>
<td>Qs/Qt (%)</td>
<td>OA 12 ± 2</td>
<td>22 ± 5*</td>
<td>21 ± 5*</td>
</tr>
<tr>
<td></td>
<td>Control 10 ± 2</td>
<td>7 ± 1</td>
<td>8 ± 1</td>
</tr>
</tbody>
</table>

Data are presented as mean ± SEM.
* Denotes a significant difference (p<0.05) time zero versus 120 min.
§ Denotes a significant difference (p<0.05) at 120 min versus 240 min within groups.
§§ Denotes significant difference (p<0.05) between groups.

Figure 1: QPORT (portal venous blood flow). Values presented as mean ± SEM (bars). For statistical comparisons see text.

Figure 2: JMP (jejunal mucosal perfusion), perfusion units (PU). Values presented as mean ± SEM (bars). For statistical comparisons see text.

Figure 3: PtiO₂ (jejunal tissue oxygen tension). Values presented as mean ± SEM (bars). For statistical comparisons see text.

Acute phase, changes between groups

Mean pulmonary arterial pressure, Qs/Qt and MO₂-ER were higher in the OA group (MO₂-ER 36 ± 1% and 28 ± 3% respectively), while pH and P/F ratio were lower in the OA group at 120 min after time zero (Table 1). Lactate flux decreased non-significantly within the OA group but was significantly lower than in the control group at 120 min after time zero (Fig. 4).

There were no significant changes in mesenteric oxygen uptake (M-VO₂) (data not shown), JMP or PtiO₂ within groups or between groups during the acute phase (Figs 2 and 3).

Late phase, changes within groups

In the OA group conditions were stable, with only slight but statistically significant decreases in MAP, CVP and MPAP (Table 1). M-DO₂ decreased from 82 ± 7 to 73 ± 6 ml min⁻¹ while MO₂-ER increased (from 36 ± 1% to 42 ± 2%) resulting in unchanged M-VO₂.

In the control group there were small but significant decreases in PaO₂, P/F ratio and PtiO₂ (PtiO₂ from 66 ± 4 to 58 ± 3 mmHg) (Table 1, Fig. 3).
Late phase, changes between groups

Lflux (mesenteric net lactate flux) and P/F ratio were lower in the OA group, while Qs/Qt and MO2-ER were higher in the OA group (MO2-ER 42 ± 2% versus 29 ± 4% respectively) when comparing groups at 240 min after time zero (Table 1, Fig. 4). There were no significant differences between groups when comparing MAP, CVP, CO, QPORT, M-DO2, M-VO2, JMP or PiO2 at 240 min after time zero.

Systemic inflammation

Systemic cytokines were measured in five animals in the OA group and six animals in the control group. TNF-α was not detected at any time in any animal. In one animal in each group, IL-6 was not detected at any time. Mean IL-6 levels in OA group were 104 ± 73, 186 ± 61 and 47 ± 27 pg ml⁻¹ at 0, 120 and 240 min respectively and 296 ± 245, 408 ± 240 and 398 ± 240 respectively. Cytokine levels were very low in all groups, and there were no significant differences between groups.

Results, ³H-OA distribution study

Haemodynamic and respiratory data (data not shown) were similar to data obtained in the OA group in the main study. Approximately 30% of the injected dose were recovered in the studied organs (lungs 16 ± 6 ± 3%, heart 1 ± 1 ± 0%, liver 9 ± 0 ± 0%, kidneys 0 ± 7 ± 0%, stomach 0 ± 3 ± 0%, jejunum 1 ± 0 ± 3, colon 0 ± 4 ± 0% and arterial blood 1 ± 5 ± 0%).

Discussion

The OA lung injury model is an established and previously well-described model, where the intestinal effects has not been clearly elucidated. The OA infusion induces a moderate to severe lung injury with hypoxemia, pulmonary capillary leakage and mechanical properties similar to those found in patients with moderate to severe ARDS (Rosenthal et al., 1998). The OA lung injury is caused both by a direct injury to the pulmonary capillary endothelium and occlusive microthrombosis (Baker et al., 1969; Derks & Jacobovitz-Derks, 1977; Grotjohan et al., 1993). It is generally considered a local lung injury model, but we have demonstrated that OA can be found in many extrapulmonary tissues 45 min after completed OA infusion. We could recover approximately 30% of the administered dose of ³H-labelled OA. The remainder was either distributed to large tissues not studied (mainly fat and muscle tissue) or already metabolized to water (Goeransson & Olivecrona, 1965). When determining ³H-OA tissue radioactivity, the contribution of residual blood ³H-OA radioactivity in the organ has to be accounted for. Using previously published data on organ residual blood volume (Hansard, 1956), we calculated corrections for each organ (data not presented), and the contribution of residual blood ³H-OA radioactivity was judged to be negligible. A small amount of ³H-labelled OA was recovered from the intestine, a total of 2% of the given dose (equal to 5% of the recovered dose). This shows that the intestines during OA infusion are exposed to a potentially inflammatory stimulus.

We measured systemic levels of the proinflammatory cytokines TNF-α and IL-6. TNF-α was not detected in any animal at any time, while IL-6 levels were generally low in both groups (in the observed range small variations in optical densities gives large variations in measured values). This finding suggests that OA infusion does not cause general systemic inflammation, but does not rule out the possibility of a local inflammatory reaction in the intestines.

When evaluating local blood flow, metabolism or oxygenation we found no indications of intestinal injury. There were no indications of intestinal hypoperfusion in either of the two groups. Mesenteric oxygen delivery and extraction ratio levels in both groups were in line with previously reported values not associated with intestinal ischaemia (Lehtipalo et al., 2001). As shown in Fig. 3, jejunal tissue oxygen tension decreased in the OA group, but only to a minor degree, and to absolute levels above ischaemic threshold values of 45 mmHg (Lehtipalo et al., 2003).

In the control group there was a constant and small net lactate production, without any signs of anaerobic conditions in the intestine. Based on in vitro data, it has been suggested that the intestine in the fed state produces lactate under aerobic conditions (Hansen & Parsons, 1976), even though this has been questioned as a possible artefact due to anaesthesia and surgical manipulation of the bowel (Kimura et al., 1988). During a metabolic challenge, such as an oral glucose load, the intestine has been shown to produce lactate under aerobic conditions in man (Bjorkman et al., 1990). These findings fit well with our notion of aerobic conditions in the control group despite a small net lactate production. In the OA group there was a large increase in arterial lactate concentrations and a corresponding slightly lesser increase in portal venous lactate concentrations. As there was a net mesenteric lactate uptake we conclude that the source of lactate production was not the intestinal vascular bed. In patients with ALI due to inflammation, net lactate production across the lung has been described by several authors (De Backer et al., 1997; Kellum et al., 1997).
In \(^{3}\mathrm{H}\)-OA-exposed animals, intestinal tissue was stained for fat and histologically examined (data not shown), but no gross evidence of inflammation such as tissue oedema or neutrophil accumulation and extravasation could be found.

A limitation of our study is the fact that animals in the OA group were given additional fluids and adrenaline during the study period in the OA group. With this knowledge at hand we conclude that the OA lung injury model is suitable for studies on intestinal effects of different treatment modalities.

Acknowledgments

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